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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|------------------|------------------------------------|----------------------|---------------------|------------------|
| 10/526,425 | 03/03/2005 | Tsuneko Okazaki | 80161(302730) | 9673 |
| | 7590 02/17/200 NGELL PALMER & D | EXAMINER | | |
| P.O. BOX 5587 | | HILL, KEVIN KAI | | |
| BOSTON, MA 02205 | | | ART UNIT | PAPER NUMBER |
| | | 1633 | | |
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| | | MAIL DATE | DELIVERY MODE | |
| | | | 02/17/2009 | PAPER |

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Advisory Action Before the Filing of an Appeal Brief

| Application No. | Applicant(s) | |
|-----------------|----------------|--|
| 10/526,425 | OKAZAKI ET AL. | |
| Examiner | Art Unit | |
| KEVIN K. HILL | 1633 | |

| | KEVIN K. HILL | 1633 |
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| The MAILING DATE of this communication appea | rs on the cover sheet with the | correspondence address |
| THE REPLY FILED <u>13 January 2009</u> FAILS TO PLACE THIS AF | PPLICATION IN CONDITION FO | OR ALLOWANCE. |
| 1. The reply was filed after a final rejection, but prior to or on the application, applicant must timely file one of the following reapplication in condition for allowance; (2) a Notice of Appear for Continued Examination (RCE) in compliance with 37 CF periods: | ne same day as filing a Notice of plies: (1) an amendment, affidav al (with appeal fee) in compliance | Appeal. To avoid abandonment of this vit, or other evidence, which places the with 37 CFR 41.31; or (3) a Request |
| a) The period for reply expires 5 months from the mailing date of b) b) The period for reply expires on: (1) the mailing date of this Advance event, however, will the statutory period for reply expire late Examiner Note: If box 1 is checked, check either box (a) or (b) MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f). | visory Action, or (2) the date set forther than SIX MONTHS from the mailing. ONLY CHECK BOX (b) WHEN TH | ng date of the final rejection. |
| Extensions of time may be obtained under 37 CFR 1.136(a). The date or have been filed is the date for purposes of determining the period of exterunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the ship set forth in (b) above, if checked. Any reply received by the Office later that may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL | nsion and the corresponding amoun ortened statutory period for reply orig | t of the fee. The appropriate extension fee ginally set in the final Office action; or (2) as |
| The Notice of Appeal was filed on A brief in compliant filing the Notice of Appeal (37 CFR 41.37(a)), or any extens Notice of Appeal has been filed, any reply must be filed with AMENDMENTS | sion thereof (37 CFR 41.37(e)), t | o avoid dismissal of the appeal. Since a |
| 3. The proposed amendment(s) filed after a final rejection, but (a) They raise new issues that would require further cons (b) They raise the issue of new matter (see NOTE below (c) They are not deemed to place the application in bette appeal; and/or | sideration and/or search (see NC); | DTE below); |
| (d) They present additional claims without canceling a continuation Sheet. (See 37 CFR 1.116 | 6 and 41.33(a)). | |
| 4. The amendments are not in compliance with 37 CFR 1.121 5. Applicant's reply has overcome the following rejection(s): _ 6. Newly proposed or amended claim(s) would be allowed. | | |
| non-allowable claim(s). 7. For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is provided the status of the claim(s) is (or will be) as follows: Claim(s) allowed: Claim(s) objected to: Claim(s) rejected: 1. 4-7. 13-14. Claim(s) withdrawn from consideration: | | rill be entered and an explanation of |
| AFFIDAVIT OR OTHER EVIDENCE | | |
| The affidavit or other evidence filed after a final action, but I because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e). | sufficient reasons why the affida | vit or other evidence is necessary and |
| 9. The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to ove showing a good and sufficient reasons why it is necessary a | ercome <u>all</u> rejections under appe | eal and/or appellant fails to provide a |
| 10. The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER | | · |
| 11. The request for reconsideration has been considered but on See Continuation Sheet. 12. Note the attached Information Disclosure Statement(s). (P | | in condition for allowance because: |
| 13. Other: | 10/00/00) 1 aper 110(5). | |
| | /Q. JANICE LI, M.D./ Primary Examiner, Art | Unit 1633 |

Continuation of 3. NOTE: The proposed amendment(s) to the claims add new limitations, thereby raising new issues which would require further search and consideration, e.g. introducing into a first mammalian host cell a first vector and introducing into a second host cell a second vector, if the proposed amendment was entered.

Continuation of 11. does NOT place the application in condition for allowance because: Claims 1, 4-6 and 13-14 stand rejected for reasons of record in the Office Action mailed August 14, 2008. Applicant requests reconsideration after final Office Action.

Claims 1, 4-6 and 13 stand rejected under 35 U.S.C. 103(a) as being unpatentable over Mejia et al in further view of Waye et al, Ikeno et al and Perkins et al.

Response to Arguments

Applicant argues that:

- a) Mejia does not teach a second vector that comprises an insertion sequence and an insulator sequence,
- b) Mejia does not teach the method being performed in mammalian cells,
- c) Perkins does not teach insertion sequences which is a loxP site or a FRT site,
- d) the combination of references fail to teach the claimed insertion sequence or insulation sequence, and
- e) the MAC of the instant invention has an insulator sequence for the purpose of promoting the expression of a gene to be introduced later, and it was found by the inventors that, surprisingly, both the efficiency of gene transfer into the mammalian artificial chromosome and the efficiency of the expression of the gene were enhanced.

Applicant's arguments have been fully considered, but are unpersuasive.

With respect to a-c), in response to applicant's arguments against the references individually, one cannot show nonobviousness by attacking references individually where the rejections are based on combinations of references. See In re Keller, 642 F.2d 413, 208 USPQ 871 (CCPA 1981); In re Merck & Co., 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986). In the instant case, Applicant appears to have overlooked that Mejia teaches a second vector being in circular form and comprising an insertion sequence for specifically inserting a sequence of interest, wherein the insertion sequence is a loxP recombination site, and that Perkins successfully demonstrated methods of producing artificial chromosomes in mammalian cells.

With respect to d), human alpha satellite insulator sequences comprising nucleic acid sequences 100% identical to SEQ ID NO:1 were known in the prior art. Thus, it is unclear how the combination of the cited references fail to teach the claimed insertion sequence (discussed immediately above) or insulation sequence.

With respect to e), as a first matter, the claims are drawn to a method of making a mammalian artificial chromosome, not a method of enhancing gene expression in a mammalian artificial chromosome. As a second matter, the insulator activities found by Applicant necessarily flow from the insulator elements, and thus would necessarily be present and achieved by the insulator elements taught in the prior art and structurally indistinguishable from the instantly claimed insulator element.

Claims 1 and 7 stand rejected under 35 U.S.C. 103(a) as being unpatentable over Mejia et al, in further view of Waye et al, Ikeno et al and Perkins et al, as applied to claims 1, 4-6 and 13 above, and in further view of Bokkelen et al. Response to Arguments

Applicant argues that Bokkelen et al do not cure the flaws of Mejia et al, Waye et al, Ikeno et al and Perkins et al.

Applicant's argument(s) has been fully considered, but is not persuasive. The Examiner's response to the arguments regarding the combination of Mejia et al, Waye et al, Ikeno et al and Perkins et al discussed above are incorporated herein. Applicant does not contest the teachings of Bokkelen et al as applied to the obviousness to substitute the mammalian centromere sequence length of Mejia et al with a mammalian centromere sequence length of about 50kb or less as taught be Bokkelen et al with a reasonable chance of success because the simple substitution of one known element for another would have yielded predictable results to one of ordinary skill in the art at the time of the invention.

Claims 1 and 14 stand rejected under 35 U.S.C. 103(a) as being unpatentable over Mejia et al, in further view of Waye et al, Ikeno et al, Perkins et al and Bokkelen et al, as applied to claims 1, 4-7 and 13 above, and in further view of Cooke et al. Response to Arguments

Applicant argues that Bokkelen et al do not cure the flaws of Mejia et al, Waye et al, Ikeno et al, Perkins et al and Bokkelen et al. Applicant's argument(s) has been fully considered, but is not persuasive. The Examiner's response to the arguments regarding the combination of Mejia et al, Waye et al, Ikeno et al, Perkins et al and Bokkelen et al discussed above are incorporated herein. Applicant does not contest the teachings of Cooke et al as applied to the obviousness to try adjusting the molecular ratio of the first and second vector in a cloning reaction to be in the range from about 10:1 to about 1:10 molecular ratio because "a person of ordinary skill has good reason to pursue the known options within his or her technical grasp. If this leads to the anticipate success, it is likely that product not of innovation but of ordinary skill and common sense." Adjusting the relative ratios between a first (donor) nucleic acid and a second (target) nucleic acid in a molecular cloning reaction has long been practiced in the art.